#### GLY-SAL 10-2- salicylic acid cloth Topiderm, Inc.

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#### Gly-Sal 10-2

#### Drug Facts

#### **Active ingredient**

Salicylic Acid USP, 2%

#### Purpose

Acne medication

### Uses

Cleansing pads for the treatment of acne, with the skin enhancement properties of Glycolic acid.

### Warnings

- For external use only.
- Keep away from eyes, lips, and mouth.
- If irritation develops, discontinue use and consult a doctor.
- Using other topical acne medication at the same time or immediately following use of this product may increase dryness or irritation of the skin. If this occurs, only one medication should be used unless directed by a doctor.
- **Keep out of reach of children.** If swallowed, seek professional assistance or contact a Poison Control Center immediately.
- Flammable; keep tightly closed, away from flame and heat.
- Sunscreen use is recommended with any Glycolic Acid product and for an additional week thereafter, because some individuals may be more sensitive to sunlight.

### Directions

- Wipe the entire affected area with a moist pad one to three times daily.
- Because excessive drying of the skin can occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a physician.
- If bothersome dryness or peeling occurs, reduce application to once a day or every other day.

### Inactive ingredients

Purified Water, Glycolic Acid, SD Alcohol 40B(13% v/v), Hamamelis Virginiana (Witch Hazel) Water, Ammonium Hydroxide, Polysorbate-20, Sodium Benzoate, Imidazolidinyl

Urea, Acetone, Disodium EDTA.

#### **PRINCIPAL DISPLAY PANEL - 60 Pad Jar**

COMPLIMENTS OF TOPIX PHARMACEUTICALS, INC

Gly / Sal 10-2 Pads

Glycolic Acid 10% Salicylic Acid USP, 2%

60 pads

Available Custom Branded 800.445.2595 c.service@topixpharm.com

#### COMPLIMENTS OF



# Gly / Sal 10-2 Pads

Glycolic Acid 10% Salicylic Acid USP, 2%

60 pads

Available **Custom Branded** 800.445.2595 c.service@topixpharm.com

Drug Facts	<b>Directions</b> Wipe the entire affected area with a moist pad one to three times
Active ingredient Purpose   Salicylic Acid USP, 2% Acne medication	daily. Because excessive drying of the skin can occur, start with one application
<b>Uses</b> Cleansing pads for the treatment of acne, with the skin enhancement properties of Glycolic acid.	daily, then gradually increase to two or three times daily if needed or as directed by a physician.
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acne medication at the same time or immediately following use of this product may increase dryness or irritation of the skin. If this occurs, only one medication should be used unless directed by a doctor. <b>Keep out of reach of children.</b> If swallowed, seek professional assistance or contact a Poison Control Center immediately. Flammable; keep tightly closed, away from flame and heat. Sunscreen use is recommended with any Glycolic Acid product and for an additional	<b>Inactive ingredients</b> Purified Water, Glycolic Acid, SD Alcohol 40B(13% v/v), Hamamelis Virginiana (Witch Hazel) Water, Ammonium Hydroxide, Polysorbate-20, Sodium Benzoate, Imidazolidinyl Urea, Acetone, Disodium EDTA.
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GLY-SAL 10-2					
salicylic acid cloth					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (So	urce)	NDC:513	26-712
Route of Administration	TOPICAL				
Active Ingredient/Active	Molety				
Ingredient Name			Basis of St	rength	Strength
SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ)			SALICYLIC ACID		20 mg
Inactive Ingredients					

Ingredient Name				Strength	
WATER (UNII: 059QF0KO0R)					
GĽ	YCOLIC ACID (UI	NII: OWT12SX38S)		100 mg	
AL	COHOL 95% (UN	II: 7528N5H79B)		130 mL	
HA	MAMELIS VIRGIN	IIANA LEAF WATER (UNII: 8FP93ED6H2)			
AM	IMONIA (UNII: 513	38Q19F1X)			
PO	LYSORBATE 20	(UNII: 7T1F30V5YH)			
so	DIUM BENZOAT	E (UNII: OJ245FE5EU)			
IMIDUREA (UNII: M629807ATL)					
AC	ETONE (UNII: 136	54PS73AF)			
ED	ETATE DISODIU	M (UNII: 7FLD91C86K)			
Packaging					
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:51326-712-	60 in 1 JAR; Type 0: Not a Combination	05/28/2019		

## **Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH DRUG	M006	05/28/2019	

## Labeler - Topiderm, Inc. (049121643)

Registrant - Topiderm, Inc. (049121643)

## Establishment

Name	Address	ID/FEI	Business Operations
Topiderm, Inc.		049121643	MANUFACTURE(51326-712)

Revised: 6/2025

Topiderm, Inc.